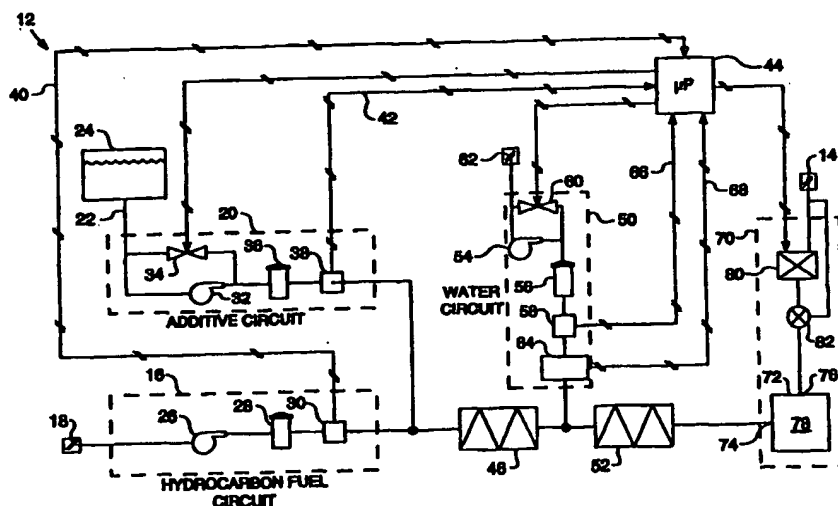


PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C10L 1/32		A1	(11) International Publication Number: WO 99/63025 (43) International Publication Date: 9 December 1999 (09.12.99)
(21) International Application Number: PCT/US99/12240 (22) International Filing Date: 2 June 1999 (02.06.99) (30) Priority Data: 60/088,068 5 June 1998 (05.06.98) US (71) Applicant: CATERPILLAR INC. [US/US]; 100 N.E. Adams Street, Peoria, IL 61629-6490 (US). (72) Inventors: COLEMAN, Gerald, N.; 2823 W. Fountaindale Drive, Peoria, IL 61614 (US). ENDICOTT, Dennis, L.; 7101 S. Stranz Road, Mapleton, IL 61547 (US). JAKUSH, Edward, A.; 2407 Park Avenue, Evanston, IL 60201 (US). NIKOLOV, Alex; Illinois Institute of Technology, 10 W. 33rd Street, Chicago, IL 60616 (US). (74) Agents: HAMPSCH, Robert, J. et al.; 100 N.E. Adams Street, Peoria, IL 61629-6490 (US).			(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>

(54) Title: **STABLE FUEL EMULSIONS AND METHOD OF MAKING**

(57) Abstract

An improved method for producing highly stable aqueous fuel emulsions. The method provides for selectively combining prescribed quantities of diesel fuel, purified water, alcohol, and an additive package and emulsifying the fuel mixture using a high shear mixer. The optimum viscosity and stability in the aqueous fuel emulsion is achieved by bypassing a selected quantity of the fuel emulsion around the high shear mixing device such that the final fuel emulsion includes a bi-modal droplet size distribution having droplet sizes less than about 15 microns in size. Additional stability and performance features of the aqueous fuel emulsion are attributed to the contents of the additive package that includes a combination of surfactants, lubricity additive, cetane improvers, citric acid, and methanol wherein each of the various additives may perform multiple functions.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroun	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

- 1 -

DescriptionSTABILE FUEL EMULSIONS AND METHOD OF MAKING5 Background of the Invention

 The present invention relates to reduced nitrogen oxide (NOx) emission fuel compositions, more particularly, to high stability aqueous fuel compositions for use in internal combustion engines.

10 Nitrogen oxides comprise a major irritant in smog and are believed to contribute to tropospheric ozone which is a known threat to health. Environmental considerations and government regulations have increased the need to reduce NOx

15 production. One problem with using diesel-fueled engines is that relatively high flame temperatures reached during combustion increase the tendency for the production of nitrogen oxides (NOx). These are formed from both the combination of nitrogen and

20 oxygen in the combustion chamber and from the oxidation of organic nitrogen species in the fuel. Various methods for reducing NOx production include the use of catalytic converters, engine timing changes, exhaust recirculation, and the burning of

25 " clean" fuels. These methods are generally too expensive and/or too complicated to be placed in widespread use. The rates at which NOx are formed is related to the flame temperature. It has been shown that a small reduction in flame temperature can result

30 in a large reduction in the production of nitrogen oxides.

- 2 -

One approach to lowering the flame temperature is to inject water in the combustion zone, however; this requires costly and complicated changes in engine design. An alternate method of using water to reduce flame temperature is the use of aqueous fuels, i.e., incorporating both water and fuel into an emulsion. Problems that may occur from long-term use of aqueous fuels include engine corrosion, engine wear, or precipitate deposition which may lead to engine problems and ultimately to engine inoperability. Problematic precipitate depositions include coalescing ionic species resulting in filter plugging and inorganic post combustion deposits resulting in turbo fouling. Another problem related to aqueous fuel compositions is that they often require substantial engine modifications, such as the addition of in-line homogenizers, thereby limiting their commercial utility.

A significant barrier to the commercial use of aqueous fuel emulsions is emulsion stability. Gravitational phase separation (during storage) and high temperature, high pressure/shear flow rate phase separation (in a working engine) of these emulsions has prevented successful commercialization.

The present invention addresses the problems associated with the use of aqueous fuel compositions by providing a stabile, inexpensive fuel emulsion with reduced NOX and particulate emissions.

30

- 3 -

Summary of the Invention

In general, the invention features a substantially ashless fuel composition that comprises hydrocarbon petroleum distillate, purified water, and an additive composition. The fuel composition preferably is in the form of an emulsion having an average droplet diameter of less than about 10 microns, which is stable at storage temperatures, as well as, at temperatures and pressures encountered during use, such as, during recirculation in a compression ignited engine.

The process for making the emulsions greatly effects the stability of the resulting compositions. The components are mixed in a serial, continuous flow process. This process allows for the continuous monitoring and instantaneous adjustment of the flow, and thus content, of each component in the final mixture. After all components are mixed, the composition is aged prior to passing it through a shear pump. The aging time is temperature dependent. The resulting emulsion is a macro-emulsion having an average droplet size of less than about 10 microns.

The amount of the hydrocarbon petroleum distillate preferably is between about 43 weight percent and about 70 weight percent of the fuel composition, more preferably between about 63 weight percent and about 68 weight percent of the fuel composition.

The amount of purified water preferably is between about 28 weight percent and about 55 weight percent of the fuel composition, more preferably between about 30 weight percent and about 35 weight

- 4 -

percent of the fuel composition. The purified water preferably contains no greater than about 50 parts per million calcium and magnesium ions, and no greater than about 20 parts per million silicon. More preferably, the purified water has a Total hardness of less than 10 parts per million and contains no greater than about 2 parts per million calcium and magnesium ions, and no greater than about 1 part per million silicon.

10 The additive composition preferably includes a surfactant and may also include one or more additives such as lubricants, corrosion inhibitors, antifreezes, ignition delay modifiers, cetane improvers, stabilizers, rheology modifiers, and the like. Individual additive ingredients may perform one or more of the aforementioned functions.

Description of Preferred Embodiments

20 Preferred fuel compositions include hydrocarbon petroleum distillates and water, preferably in the form of an emulsion. The preferred emulsion is a stable system with as little surfactant as possible. A stable emulsion is desirable because a separate water phase will lead to combustion problems. Stability means no substantial phase separation in long term storage under typical storage conditions, for example up to about three months. A small amount of phase separation in the storage tank containing the fuel composition may be tolerated because pumping the fuel composition will ensure sufficient emulsification. High temperature, high pressure

- 5 -

stability is also required to maintain the emulsion under operating conditions.

The fuel composition is preferably ashless. For the purposes of this disclosure "ashless" means that, once the fuel components are combined, the level of particulates and coalescing ionic species is sufficiently low to allow long-term operation of the internal combustion engine (for example, substantially continuous operation for three months) without significant particulate and coalescing ionic species deposition on engine parts, including valve seats and stems, injectors and plug filters, and post-combustion engine parts such as the exhaust trains and turbo recovery units. The level of ash is determined by monitoring water purity, exhaust emissions, and by engine autopsy. Engine autopsy, including dismantlement and metallurgical analysis, is also used to analyze corrosion and wear.

Preferred compositions include about 43% to about 70% by weight hydrocarbon petroleum distillate, more preferably about 63% to about 68% hydrocarbon petroleum distillate. The amount and type of hydrocarbon petroleum distillate is selected so that the kilowattage per gallon provided by combusting the fuel composition is sufficiently high so that the engine need not be derated. Such hydrocarbon petroleum distillates include high paraffinic, low aromatic hydrocarbon petroleum distillates having an aromatic content of less than about 10%, preferably less than about 3%. Examples of suitable hydrocarbon petroleum distillates include kerosene, diesel, naphtha, and aliphatics and paraffinics, used alone or

- 6 -

in combination with each other, with diesel being preferred, for example, EPA Emissions Certification Diesel Fuel and standard number 2 diesel.

5 The water component of the fuel composition functions to reduce NOx and particulate emissions. The greater the amount of water, the greater the decrease in NOx emissions. The current upper limit of water is about 55%, above which the burning characteristics of the fuel make it's use impractical
10 under normal conditions, i.e., with an acceptable amount of additives and relatively inexpensive hydrocarbon petroleum distillate. The preferred amount of purified water is between about 28 weight percent and about 55 weight percent of the fuel composition,
15 more preferably between about 30 weight percent and about 35 weight percent of the fuel composition.

 The water is preferably purified such that it contains very low concentrations of ions and other impurities, particularly calcium ions, magnesium ions,
20 and silicon. This is desirable because impure water contributes to ashing and engine deposit problems after long-term use, which can lead to wear, corrosion, and engine failure. The purified water preferably contains no greater than about 50 parts per
25 million calcium and magnesium ions, and no greater than about 20 parts per million silicon. More preferably, the purified water has a Total hardness of less than 10 parts per million and contains no greater than about 2 parts per million calcium and magnesium
30 ions, and no greater than about 1 part per million silicon. Suitable purification techniques are well-known and include distillation, ion exchange

- 7 -

treatment, and reverse osmosis, with reverse osmosis being preferred due to lower cost and ease of operation.

5 The composition preferably includes an additive containing, for example, one or more of the following: surfactants, lubricants, corrosion inhibitors, antifreezes, ignition delay modifiers, cetane improvers, stabilizers, rheology modifiers, and the like. The amount of additive selected is
10 preferably sufficiently high to perform its intended function and, preferably sufficiently low to control the fuel composition cost. The additives are preferably selected so that the fuel composition is ashless.

15 In a preferred embodiment the water functions as the continuous phase of an emulsion, acting as an extender and carrier of the hydrocarbon petroleum distillate. As the continuous phase, the lower useable limit of water is theoretically about
20 26%, below which point the physics of the system inhibits maintaining water as the continuous phase.

 The composition preferably includes surfactant which facilitates the formation of a stable emulsion of the hydrocarbon petroleum distillate
25 within the continuous water phase. A preferred surfactant is a surfactant package comprised of one or more surfactants in combination with one or more surfactant stabilizers. Preferred surfactants are ashless and do not chemically react with other
30 components in the fuel composition. Suitable surfactants include nonionic, anionic and amphoteric surfactants. Preferred fuel compositions include about

- 8 -

0.3% to about 1.0% by weight, preferably about 0.4% to about 0.6% total surfactant. Examples of surfactants include alkylphenolethoxylates, alcohol ethoxylates, fatty alcohol ethoxylates, and alkyl amine ethoxylates. Of these, the alkylphenolethoxylates and alcohol ethoxylates are preferred. Of the alkylphenolethoxylates, polyethoxylated nonylphenols having between 8 and 12 moles of ethylene oxide per mole of nonylphenol are preferred. An example nonylphenol, 2,6,8-Trimethyl-4-nonyloxypolyethyleneoxyethanol is commercially available, e.g., from Union Carbide under the trade designation "TERGITOL TMN-10". Another nonylphenol ethoxylate NP-9 available from Shell under the trade designation "NP-9EO", added at 1000-3000 ppm. A preferred alcohol ethoxylate is a C₁₁ alcohol ethoxylate with 5 moles of ethylene oxide per mole of alcohol commercially available from Shell as "Neodol N1-5 Surfactant". Additional preferred surfactant components include, for example, Pluronic 17R-2 [octylphenoxypolyethoxyethanol] (a block copolymer produced by BASF) added at 100 - 300 ppm; CA-720 an octylphenol aromatic ethoxylate available from Rhone-Poulenc as "Igepal CA-720" added at 1000-3000 ppm; and X-102 an ethoxylated alkyl phenol available from Union Carbide as "TRITON X-102" added at 1000 - 2000 ppm.

The fuel composition preferably includes one or more lubricants to improve the slip of the water phase and for continued smooth operation of the fuel delivery system. The amount of lubricant generally ranges from about 0.04% to 0.1% by weight, more

- 9 -

preferably from 0.04% to 0.05% by weight. Suitable lubricants include a combination of mono-, di-, and tri-acids of the phosphoric or carboxylic types, adducted to an organic backbone. The carboxylic types are more preferred because of their ashless character. Examples include mixed esters of alkoxyated surfactants in the phosphate form, and di- and tri-acids of the Diels-Alder adducts of unsaturated fatty acids. The organic backbone preferably contains about 12 to 22 carbons. A specific example of a suitable lubricant is Diacid 1550™ (Atrachem Latol 1550 or Westvaco Chemicals Diacid 1550), which is preferred due to its high functionality at low concentrations. The Diacid 1550 also has nonionic surfactant properties.

Neutralization of the phosphoric and carboxylic acids, preferably with an alkanolamine, reduces possible corrosion problems caused as a result of the addition of the acid. Suitable alkanolamine neutralizers include amino methyl propanol, triethanolamine, and diethanolamine, with amino methyl propanol (available from Angus Chemical under the trade designation "AMP-95") being preferred. Preferred compositions include about 0.05 to 0.4% by weight neutralizer, more preferably about 0.06%.

The fuel composition may also include one or more corrosion inhibitors, preferably one that does not contribute a significant level of inorganic ash to the composition. Aminoalkanoic acids are preferred. An example of a suitable corrosion inhibitor is available from the Keil Chemical Division of Ferro Corporation under the trade designation "Synkad 828".

- 10 -

Preferred compositions include about 0.05% by weight corrosion inhibitor.

The fuel composition may also include one or more ignition delay modifiers, preferably a cetane improver, to improve fuel detonation characteristics, particularly where the fuel composition is used in compression ignited engines. Examples include nitrates, nitrites, and peroxides. The preferred ignition delay modifier is 2-ethylhexylnitrate (2-EHN), available from Ethyl Corporation under the trade designation "HiTec 4103". Ammonium nitrate can also be used as a known cetane improver. Preferred compositions include about 0.1% to 0.4% by weight ignition delay modifier.

An antifreeze may also be included in the fuel composition. Organic alcohols are preferred. Specific examples include methanol, ethanol, isopropanol, and glycols, with methanol being preferred. The amount of antifreeze preferably ranges from about 2% to about 9% by weight.

Biocides known to those skilled in the art may also be added, provided they are ashless. Antifoam agents known to those skilled in the art may be added as well, provided they are ashless. The amount of antifoam agent preferably is not more than .0005% by weight.

The fuel composition may also include one or more coupling agents (hydrotropes) to maintain phase stability at high temperatures and shear pressures. High temperature and shear pressure stability is required, for example, in compression ignited (diesel) engines because all the fuel delivered to the

- 11 -

injectors may not be burned to obtain the required power load in a given cycle. Thus, some fuel may be recirculated back to the fuel tank. The relatively high temperature of the recirculated fuel, coupled with the shear pressures encountered during recirculation, tends to cause phase separation in the absence of the coupling agent.

Examples of preferred coupling agents include di- and tri-acids of the Diels-Alder adducts of unsaturated fatty acids. A specific example of a suitable coupling agent is Diacid 1550, neutralized with an alkanolamine to form a water soluble salt. Suitable alkanolamine neutralizers include amino methyl propanol, triethanolamine, and diethanolamine, with amino methyl propanol preferred. The amount of the coupling agent typically ranges from about 0.04% to 0.1 % by weight, more preferably 0.04 to 0.05%.

The fuel composition includes additives which perform multiple functions. For example, Diacid 1550 acts as a surfactant, lubricant, and coupling agent. Similarly, AMP-95 acts as a neutralizer and helps maintain the pH of the fuel composition and, if used, ammonium nitrate acts as a cetane improver and an emulsion stabilizer..

A preferred fuel composition has the following composition: 67% by weight diesel, 30% by weight water, 2% by weight methanol, 0.16% by weight X-102; 0.08% by weight N1-5; 0.08% by weight TMN-10, 0.04% Diacid 1550, 0.06% AMP-95, 0.05% Synkad 828, and 0.37% 2-ethylhexylnitrate.

- 12 -

Emulsion Process

The fuel compositions are manufactured using a continuous flow process capable of continuous monitoring and adjustment which can provide the shear rates necessary to form the desired droplet size for a stable emulsion.

The process uses a fuel emulsion blending system including a first inlet circuit adapted for receiving hydrocarbon petroleum distillate from the source of hydrocarbon petroleum distillate; a second inlet circuit adapted for receiving aqueous fuel emulsion additives from the source of aqueous fuel emulsion additives; a third inlet circuit adapted for receiving water from the source of water. The blending system further includes a first blending station adapted to mix the hydrocarbon petroleum distillate and aqueous fuel emulsion additives and a second blending station adapted to mix the hydrocarbon and additive mixture received from the first blending station together with the water received from the source of water. The blending system further includes an emulsification station downstream of the blending stations, which is adapted to emulsify the mixture of hydrocarbon petroleum distillate, additives and water to yield a stable aqueous fuel emulsion. The present embodiment of the blending system is operatively associated with a blending system controller which is adapted to govern the flow of the hydrocarbon petroleum distillate, water and aqueous fuel emulsion additives thereby controlling the mixing ratio in accordance with prescribed blending ratios.

- 13 -

In an example of a continuous process, the additives are combined in the form of a stream, and then fed to a first in-line blending station where they are combined with a hydrocarbon petroleum distillate stream. The resulting product is then combined with water in a second in-line blending station to form the fuel composition, which is then aged in a reservoir and then pumped using a shear pump to a storage tank. In an alternate embodiment, a separate stream of the antifreeze (alcohol) is combined with the other additives in an in-line blending station and then this combined additive stream is fed to the first in-line blending station.

Figure 1 illustrates a schematic representation of a preferred aqueous fuel emulsion blending system 12 having a plurality of ingredient inlets and an aqueous fuel emulsion outlet 14. As seen therein, the preferred embodiment of the fuel blending system 12 comprises a first fluid circuit 16 adapted for receiving hydrocarbon petroleum distillate at a first ingredient inlet 18 from a source of hydrocarbon petroleum distillate (not shown) and a second fluid circuit 20 adapted for receiving fuel emulsion additives at a second ingredient inlet 22 from an additive storage tank 24 or similar such source of fuel emulsion additives. The first fluid circuit 16 includes a fuel pump 26 for transferring the hydrocarbon petroleum distillate, preferably a diesel fuel, from the source of hydrocarbon petroleum distillate to the blending system 12 at a selected flow rate, a 10 micron filter 28, and a flow

- 14 -

measurement device 30 adapted to measure the flow rate of the incoming hydrocarbon petroleum distillate stream. The second fluid circuit 22 also includes a pump 32 for transferring the additives from the storage tank 24 to the blending system 12 at prescribed flow rates. The fuel additive flow rate within the second fluid circuit 20 is controlled by a flow control valve 34 interposed between the additive storage tank 24 and the pump 26. As with the first fluid circuit 16, the second fluid circuit 20 also includes a 10 micron filter 36 and a flow measurement device 38 adapted to measure the controlled flow rate of the incoming additive stream. The signals 40, 42 generated from the flow measurement devices 30, 38 associated with the first and second fluid circuits are further coupled as inputs to a blending system controller 44.

The first fluid circuit 16 transporting the hydrocarbon petroleum distillate and the second fluid circuit 20 adapted for supplying the fuel additives are coupled together and subsequently mixed together using a first in-line mixer 46. The resulting mixture of hydrocarbon petroleum distillate and fuel additives is then joined with a purified water stream supplied via a third fluid circuit 50 and subsequently mixed together using a second in-line mixer 51.

The third fluid circuit 50 includes a water pump 54 for transferring the purified water from a source of clean or purified water (not shown) at a selected flow rate to the blending system 12, a particulate filter 56 and a flow measurement device 58 adapted to measure the flow rate of the incoming

- 15 -

purified water stream. The water pump 54, filter 56 and flow measurement device 58 are serially arranged within the third fluid circuit 50. The water flow rate within the third fluid circuit 50 is preferably controlled using a flow control valve 60 interposed between the clean water source and the water pump 54 proximate the third or water inlet 62. The third fluid circuit 50 also includes a specific conductance measurement device 64 disposed downstream of the flow measurement device 58 and adapted to monitor the quality of the water supplied to the blending system 12. The signals 66, 68 generated from the flow measurement device 58 and the specific conductance measurement device 64 in the third fluid circuit 50 are provided as inputs to the blending system controller 44. If the water quality is too poor or below a prescribed threshold, the blending system controller 44 disables the blending system 12 until corrective measures are taken. In the preferred embodiment, the water quality threshold, as measured using the specific conductance measurement device 64 should be no greater than 20 microsiemens per centimeter. As indicated above, the purified water from the third fluid circuit 50 is joined with the hydrocarbon petroleum distillate and fuel additive mixture and subsequently re-mixed using the second in-line mixer 52 or equivalent blending station equipment.

The resulting mixture or combination of hydrocarbon petroleum distillate, fuel emulsion additives, and purified water are fed into an emulsification station 70. The emulsification station

- 16 -

70 includes an aging reservoir 72, and emulsifier. The aging reservoir 72 includes an inlet 74, an outlet 76 and a high volume chamber 78 or reservoir. The preferred embodiment of the blending system 12 operates using a three-minute aging time for the aqueous fuel emulsion. In other words, a blending system operating at an output flow rate of about 15 gallons per minute would utilize a 45-gallon tank as an aging reservoir. The incoming stream of hydrocarbon petroleum distillate, fuel emulsion additives, and purified water are fed into the aging reservoir 72 at a location that preferably provides continuous agitation to the reservoir. The preferred embodiment of the blending system 12 also includes a high shear pump 80 and a pressure regulating valve 32 disposed downstream of the aging reservoir 72 which provides the final aqueous fuel emulsion at the blending system outlet 14.

As indicated above, the blending system controller 44 accepts as inputs the signals generated by the various flow measurement devices in the first, second and third fluid circuits, as well as any signals generated by the water quality measurement device together with various operator inputs such as prescribe fuel mix ratios and provides control signals for the flow control valve in the second fluid circuit and the flow control valve in the third fluid circuit. The illustrated embodiment of the blending system is preferably configured such that the hydrocarbon petroleum distillate stream is not precisely controlled by is precisely measured. Conversely, the purified water feed line and the fuel additive feed

- 17 -

line are precisely controlled and precisely measured to yield a prescribed water blend fuel mix. The illustrated embodiment also shows the hydrocarbon petroleum distillate, purified water and fuel additive streams to be continuous feed so that the proper fuel blend ratio is continuously delivered to the shear pump. Alternatively, however, it may be desirable to configure the blending system such that the purified water stream is precisely measured but not precisely controlled while precisely controlling and measure the hydrocarbon petroleum distillate feed line and the fuel additive feed line to yield a prescribed water blend fuel mix.

Examples of shear pumps capable of the necessary high shear rates are the Ross X Series mixer and the Kady Mill. The preferred shear pump is a Kady Mill, running at between about 20 Hz and 60 Hz, preferably about 40Hz. As in the case of the batch process, the product is in the form of a stable, homogeneous, milky emulsion having an average droplet diameter of less than 10 microns, preferably ranging from about 4 to about 6 microns.

In another preferred embodiment of the process, 10% to 50% of the flow from the reservoir is sent directly to the storage tank, by-passing the shear pump. This results in an emulsion having bi-modal distribution of preferred droplet diameters. The bi-modal distribution enhances stability while allowing for an optimal viscosity. The percentage is adjusted to obtain optimal viscosity and stability of the final emulsion.

Engine Design

The aqueous fuel compositions according to the invention can be used in internal combustion engines without substantially modifying the engine design. For example, the fuel compositions can be used without re-designing the engine to include in-line homogenizers. To enhance fuel efficacy, however, several readily implemented changes are preferably incorporated in the engine structure.

The capacity of the engine fuel system may be increased to use the fuel compositions in diesel engines. The increased capacity is a function of the percentage of water in the fuel. The engine fuel system capacity is typically scaled by the following ratio:

$$\frac{\text{Lower Heating Value of Diesel Fuel (btu/gal)}}{\text{Lower Heating Value of Fuel Composition (btu/gal)}}$$

In many cases, the engine fuel system capacity can be increased sufficiently by increasing the injector orifice size. Other engines may require an increase in the capacity of the injection pump. In addition, an increase in the capacity of the fuel transfer pump may be required.

Some modifications to the engine may be required to compensate for fuel compositions with cetane quality lower than diesel fuel. This may include advancing the fuel injection timing to improve operation at light load, during starting, and under warm up conditions. In addition, a jacket water

aftercooler may be required to warm the intake air under light load conditions. The use of a block heater or an inlet air heater may be required to improve cold starting capability.

- 5 The following examples will further describe the invention. These examples are intended only to be illustrative. Other variations and modifications may be made in form and detail described herein without departing from or limiting the scope of the invention
10 which is determined by the attached claims.

EXAMPLES DO NOT INCLUDE AN EXAMPLE OF
CONTINUOUS FLOW PROCESS WE ARE CLAIMING!!!

15 EXAMPLE 1

 An aqueous fuel composition having the following formula was prepared:

	ppm	Percent
Diesel fuel	(balance)	67%
H2O	300,000	30.00%
MeOH	20,000	2.00%
X-102	1,600	0.16%
N1-5	800	0.08%
TMN-10	800	0.08%
DA-1550	400	0.04%
AMP-95	600	0.06%
Synkad 828	500	0.05%
2-EHN	3,700	0.37%

- 20 -

The fuel is prepared by first mixing the Diacid 1550, AMP-95, Synkad 828, X-102, N1-5, and TMN-10 with the methanol. The mixture is agitated.

5 The mixture is charged into a vessel with the reverse osmosis purified water and agitated for about 1-5 minutes. Then the F-173 and 2-ethylhexyl nitrate are charged into the vessel, and the composition is agitated for 15-30 minutes. The mixing vessel is a Lightnin Blender, and all mixing is carried out under
10 ambient conditions.

The aqueous fuel composition is then pumped through a Kady Mill shear pump at a shear rate of 40 Hz resulting in a homogeneous, milky emulsion having an average droplet diameter of about 4 to about 6
15 microns. The fuel composition is stored at ambient temperatures.

EXAMPLE 2

20 A fuel composition was prepared by the method of Example 1, having the formula:

Diesel Fuel	67%
Highly purified water	30%
Methanol	2.00%
2-EHN	0.37%
DA-1550	400 ppm
AMP 95	600 ppm
Synkad 828	500

- 21 -

	ppm
N1-5	1000 ppm
NP 9	3000 ppm

EXAMPLE 3

5 A fuel composition was prepared by the method of Example 1, having the formula:

Diesel Fuel	67%
Highly purified water	30%
Methanol	2.00%
2-EHN	0.37%
DA-1550	400 ppm
AMP 95	600 ppm
Synkad 828	500 ppm
TMN 10	1000 ppm
NP 9	2000 ppm
17R2	100 ppm

EXAMPLE 4

10 A fuel composition was prepared by the method of Example 1, having the formula:

Diesel Fuel	67%
Highly purified water	30%
Methanol	2.00%

- 22 -

2-EHN	0.37%
DA-1550	400 ppm
AMP 95	600 ppm
Synkad 828	500 ppm
N1-5	1000 ppm
TMN 10	1000 ppm
CA 720	2000 ppm

For Examples 1 - 4:

the Diesel fuel used was EPA Emissions
Certification Diesel Fuel;

5 the water was purified by reverse osmosis;
X-102 is Union Carbide Triton X-102;
TMN-10 is Union Carbide Tergitol TMN-10
surfactant;

N1-5 is Shell Neodol N1-5 surfactant;

10 DA-1550 is Atrachem Latol 1550 (or Westavco
Chemicals Diacid 1550);

AMP-95 is 2-amino-2-methyl-1-propanol.

Synkad 828 is Ferro Synkad 828;

2-EHN is Ethyl Corp. 2-ethylhexyl nitrate;

15 CA-720 is Rhone-Poulenc "Igepal CA-720" ;

NP 9 is Shell "NP-9EO" ;; and

17R2 is BASF "Pluronic 17R-2" .

Example 5

20 The fuel compositions prepared according to
Examples 1, 2, 3, and 4 were run in a diesel engine to

monitor NOx and particulate emissions. The engine used was a Caterpillar 12 liter compression-ignited truck engine (four stroke, fully electronic, direct injected engine with electronic unit injectors, a turbocharger, and a four valve quiescent head) The Caterpillar C-12 truck engine was rated at 410 hp at 1800 rpm with a peak torque of 2200 N-m at 1200 rpm and was modified to run a fuel-in-water emulsion. A simulated air-to-air aftercooler (43°C inlet manifold temperature) was used.

The electronic unit injectors were changed to increase the quantity of fuel injected into the cylinder. As modified, the electronic unit injector Caterpillar Part Number 116-8800 replaced the standard injector Caterpillar Part Number 116-8888. In addition, the electronic control strategy was optimized with respect to emissions, fuel consumption, and cold starting.

Tests were performed on standard diesel fuels and on fuel emulsions of Example 1 and fuel emulsions prepared as in Example 1 in which the diesel fuel was Carb Diesel; RME (Rapeseed Methyl Ester); and Fischer Tropsch diesel. The tests were performed at 1800 rpm and 228 kW, 122 rpm and 197 kW, and 1800 rpm and 152 kW. Particulate emissions and NOx+HC emissions for standard diesel fuels and for fuel emulsions are shown in the following table:

Engine		Standard diesel fuel	Fuel emulsions
1800 rpm 228 kW	Particulate emissions (g/hp-hr)	about 0.040 to about 0.055	about 0.025 to about 0.055

- 24 -

	NOx + HC emissions (g/hp-hr)	about 2.5 to about 4.5	about 1.0 to about 2.8
1200 rpm 197 kW	Particulate emissions (g/hp-hr)	about 0.03 to about 0.033	about 0.028 to about 0.1
	NOx + HC emissions (g/hp-hr)	about 3.5 to about 6.5	about 1.5 to about 4.2
1800 rpm 152 kW	Particulate emissions (g/hp-hr)	about 0.068 to about 0.084	about 0.038 to about 0.050
	NOx + HC emissions (g/hp-hr)	about 2.3 to about 4.5	about 1.1 to about 2.7

Example 6

5 The Ball on Three Disks (BOTD) lubricity test was
 utilized to assess the lubricity of the fuel
 compositions. This test was developed by Falex
 Corporation to assess the lubricity of various diesel
 fuels and their additives. The average wear scar
 diameter is used to assess fuel composition lubricity;
 10 a smaller scar diameter implies a higher fuel
 composition lubricity. Typical diesel fuel will have
 a scar diameter of 0.45mm to 0.55mm. Fuel emulsions of
 Example 1 and fuel emulsions prepared as in Example 1
 in which the diesel fuel was Carb Diesel; RME(Rapeseed
 15 Methyl Ester); and Fischer Tropsch diesel, ranged from
 about 0.547 to about 0.738.

- 25 -

Claims

1. A high stability, low emission, fuel emulsion composition for an internal combustion engine comprising: purified water; hydrocarbon petroleum distillate; and additive; said emulsion being made by a continuous flow process comprising the steps of:

a) blending a flow of said additive and a flow of said hydrocarbon petroleum distillate in an in-line blending station;

b) blending a flow from the in-line blending station of step a) with a flow of said purified water in a second in-line blending station;

c) aging the composition from the second in-line blending station of step b) in a reservoir;

d) passing the aged composition from step c) through a shear pump to a storage tank; said emulsion having an average droplet size of less than about 10 microns.

2. The fuel emulsion composition of claim 1 comprising 26-55% purified water.

3. The fuel emulsion composition of claim 1 comprising 43-70% hydrocarbon petroleum distillate.

4. The fuel emulsion composition of claim 1 comprising less than 10% additive.

5. The fuel emulsion composition of claim 1 wherein said additive comprises comprising:

- 26 -

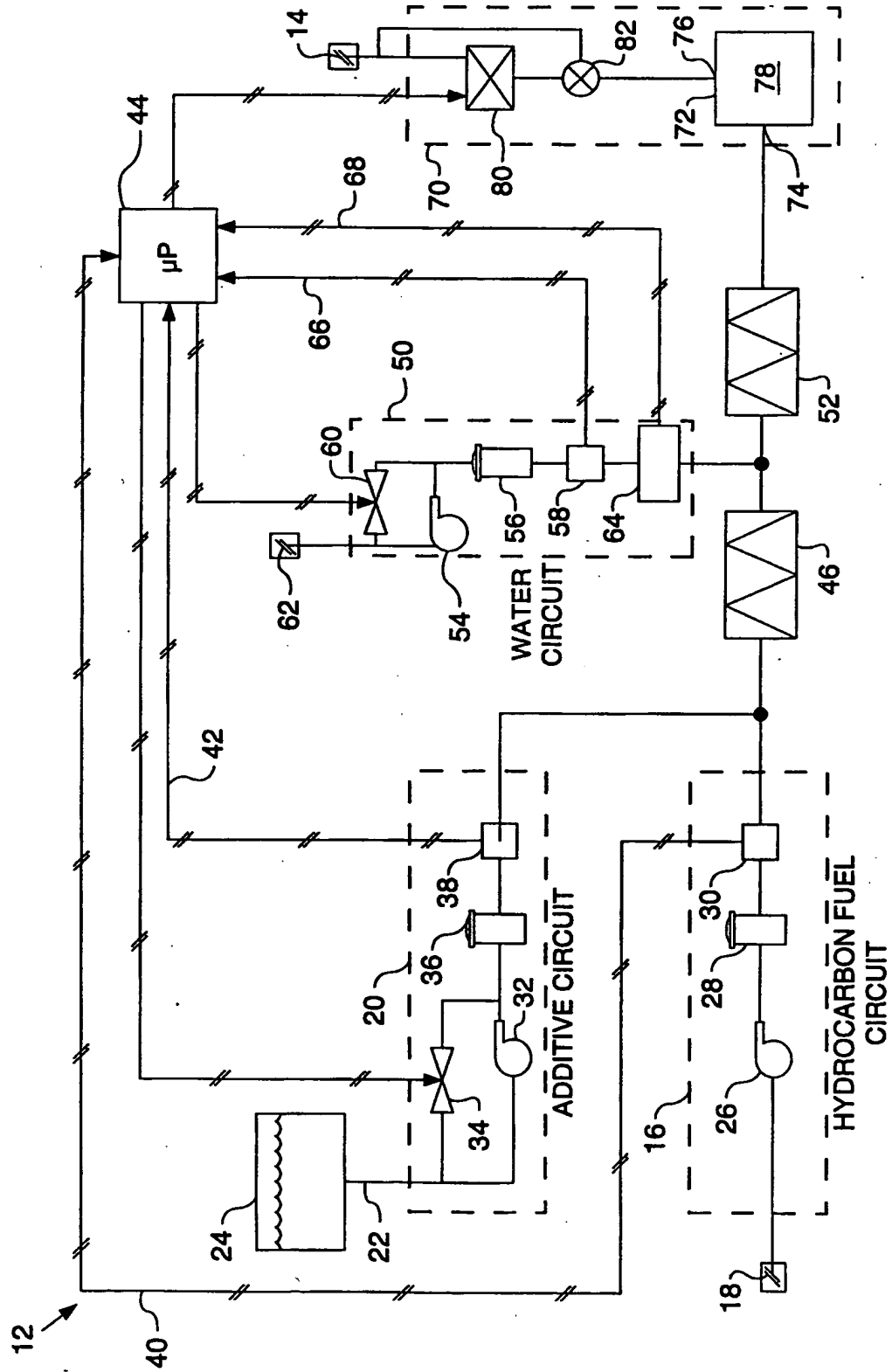
surfactant; lubricant; corrosion inhibitor; anti-freeze; and ignition delay modifier.

5 6. The fuel emulsion composition of claim
1 comprising diesel fuel, purified water, methanol,
X-102; N1-5; TMN-10, Diacid 1550, AMP-95, Synkad 828,
and 2-ethylhexylnitrate.

10 7. The fuel emulsion composition of claim
6 comprising 67% by weight diesel fuel, 30% by weight
purified water, 2% by weight methanol, 0.16% by weight
X-102; 0.08% by weight N1-5; 0.08% by weight TMN-10,
0.04% Diacid 1550, 0.06% AMP-95, 0.05% Synkad 828, and
0.37% 2-ethylhexylnitrate.

15 8. The fuel emulsion composition of claim
1 wherein said flow of additives of step a) is
comprised of a blended flow of a flow of antifreeze
and a flow of said remaining additives blended in
20 another in-line blending station.

 9. The fuel emulsion composition of claim
1 wherein in step d) only 50% to 90% of the aged
composition from step c) is passed through a shear
25 pump to a storage tank and the remaining 10% to 50% is
passed directly into the storage tank.



INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/12240

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C10L1/32

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C10L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 95 27021 A (GUNNERMAN RUDOLF W) 12 October 1995 (1995-10-12) claims 1,3,19,26	1,3,4,6, 7
P,X A	US 5 873 916 A (SCHEUERMANN TED W ET AL) 23 February 1999 (1999-02-23) claims 1,2 column 3, line 63 - column 4, line 35	1-5,8,9. 1,2
A	EP 0 301 766 A (BRITISH PETROLEUM CO PLC) 1 February 1989 (1989-02-01) claims 7,9,18 page 4, line 4	1-3

-/-



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"A" document member of the same patent family

Date of the actual completion of the international search

13 September 1999

Date of mailing of the international search report

24/09/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

De Herdt, O

INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/US 99/12240

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>WO 98 12285 A (BERTHA ANDRAS ;FUELOEP LEVENTE (HU)) 26 March 1998 (1998-03-26) claim 1 example 2</p>	1-5

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/12240

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9527021 A	12-10-1995	AU 687189 B AU 2232495 A BG 100888 A BR 9507273 A CA 2187076 A CN 1147830 A CZ 9602916 A EP 0754214 A FI 963957 A HU 76441 A JP 9511540 T NO 964163 A NZ 283877 A PL 316690 A SK 126296 A ZA 9502753 A	19-02-1998 23-10-1995 31-07-1997 23-09-1997 12-10-1995 16-04-1997 12-03-1997 22-01-1997 03-12-1996 28-08-1997 18-11-1997 04-12-1996 22-09-1997 03-02-1997 09-07-1997 21-12-1995
US 5873916 A	23-02-1999	WO 9941339 A	19-08-1999
EP 0301766 A	01-02-1989	AU 609501 B AU 2000188 A DE 3879309 A JP 1048894 A NO 174330 B SU 1793953 A US 5000757 A	02-05-1991 02-02-1989 22-04-1993 23-02-1989 10-01-1994 07-02-1993 19-03-1991
WO 9812285 A	26-03-1998	HU 9602590 A AU 3043797 A	28-10-1998 14-04-1998